

University of Groningen

Cerebral blood flow velocity changes during upright positioning in bed after acute stroke

Aries, Marcel J; Elting, Jan Willem; Stewart, Roy; De Keyser, Jacques; Kremer, Berry; Vroomen, Patrick

Published in:
BMJ Open

DOI:
[10.1136/bmjopen-2013-002960](https://doi.org/10.1136/bmjopen-2013-002960)

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version
Publisher's PDF, also known as Version of record

Publication date:
2013

[Link to publication in University of Groningen/UMCG research database](#)

Citation for published version (APA):

Aries, M. J., Elting, J. W., Stewart, R., De Keyser, J., Kremer, B., & Vroomen, P. (2013). Cerebral blood flow velocity changes during upright positioning in bed after acute stroke: An observational study. *BMJ Open*, 3, [e002960]. <https://doi.org/10.1136/bmjopen-2013-002960>

Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

The publication may also be distributed here under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license. More information can be found on the University of Groningen website: <https://www.rug.nl/library/open-access/self-archiving-pure/taverne-amendment>.

Take-down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): <http://www.rug.nl/research/portal>. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.

Cerebral blood flow velocity changes during upright positioning in bed after acute stroke: an observational study

Marcel J Aries,¹ Jan Willem Elting,¹ Roy Stewart,² Jacques De Keyser,^{1,3} Berry Kremer,¹ Patrick Vroomen¹

To cite: Aries MJ, Elting JW, Stewart R, *et al.* Cerebral blood flow velocity changes during upright positioning in bed after acute stroke: an observational study. *BMJ Open* 2013;**3**:e002960. doi:10.1136/bmjopen-2013-002960

► Prepublication history and additional material for this paper is available online. To view these files please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2013-002960>).

Received 27 March 2013

Revised 17 July 2013

Accepted 19 July 2013

ABSTRACT

Objectives: National guidelines recommend mobilisation in bed as early as possible after acute stroke. Little is known about the influence of upright positioning on real-time cerebral flow variables in patients with stroke. We aimed to assess whether cerebral blood flow velocity (CBFV) changes significantly after upright positioning in bed in the acute stroke phase.

Design: Observational study.

Participants: 47 patients with acute ischaemic stroke measured in the subacute phase after symptom onset and 20 healthy controls.

Primary and secondary outcome measures: We recorded postural changes in bilateral transcranial Doppler (primary outcome) and simultaneously recorded near-infrared spectroscopy, end-tidal CO₂, non-invasive blood pressure data and changes in neurological status (secondary outcomes).

Methods: Postures included the supine, half sitting (45°), sitting (70°) and Trendelenburg (−15°) positions. Using multilevel analyses, we compared postural changes between hemispheres, outcome groups (using modified Rankin Scale) as well as between patients and healthy controls.

Results: The mean patient age was 62±15 years and median National Institute of Health Stroke Scale score on admission was 7 (IQR 5–14). Mean proportional CBFV changes on sitting were not significantly different between healthy controls and affected hemispheres in patients with stroke. No significant differences were found between affected and unaffected stroke hemispheres and between patients with unfavourable and favourable outcomes. During upright positioning, no neurological worsening or improvement was observed in any of the patients.

Conclusions: No indications were found that upright positioning in bed in mild to moderately affected patients with stroke compromises flow and (frontal) oxygenation significantly during the subacute phase of stroke. Supine or Trendelenburg positioning does not seem to augment real-time flow variables.

ARTICLE SUMMARY

Article focus

- Stroke units save more patients with stroke from death or dependency than any other intervention in stroke.
- Early upright positioning is an intervention in the stroke unit that is recommended in several national guidelines on the basis of presumed benefits, though evidence is lacking.
- This study aimed to assess whether cerebral blood flow velocity (CBFV) —as a surrogate marker for cerebral perfusion—changes significantly after upright positioning in bed in the subacute stroke phase.

Key messages

- Following acute ischaemic stroke in the middle cerebral artery territory, early upright positioning in bed resulted in minor average changes in CBFV and cerebral oxygenation only and therefore appears to be safe during the subacute phase on the stroke unit.
- Our work is the largest observational study about real-time postural haemodynamic changes so far and is important in further unravelling the benefits of stroke unit care and early mobilisation in particular.

Strengths and limitations of this study

- The study is an example of how a small-scale study of an important surrogate endpoint can help define stroke unit management when evidence from large-scale clinical trials is not yet available. We feel a wider application of such bedside techniques in similar studies in the stroke unit is also an important means to increase the gain of subsequent larger clinical trials.
- Replication of this study should be considered when studying the (hyper)acute stroke phase, longer sitting periods, active standing and using more sophisticated and direct methods that measure cerebral perfusion regionally and estimate cerebral oxygen metabolism.

¹Department of Neurology, University of Groningen, University Medical Center Groningen, Groningen, The Netherlands

²Health Sciences, Community and Occupational Medicine, University of Groningen, University Medical Center Groningen, Groningen, The Netherlands

³Center for Neurosciences, Vrije Universiteit Brussel (VUB), Brussels, Belgium

Correspondence to

Dr Marcel Aries;
m.j.h.aries@umcg.nl

INTRODUCTION

Brain ischaemia is a potentially reversible process that is dependent on restoration of

cerebral blood flow (CBF) within a time window of cellular viability that varies according to the severity and duration of flow

cessation.¹ Any activity that alters CBF during the acute phase of ischaemic stroke may, at least theoretically, directly contribute to extension of the penumbra to the ischaemic core and compromise clinical outcome. National guidelines recommend mobilisation in and out of bed as early as possible after stroke.^{2–4} Such early upright positioning prevents choking, cardiopulmonary deconditioning and may improve early plasticity of the brain.^{5–6} However, there is insufficient evidence in support of early upright positioning after acute stroke.⁷ Also, there is fear that mobilisation in the setting of a (potentially) disturbed cerebral autoregulation leads to gravity induced diminished CBF to the penumbra and, potentially, extension of the infarct core.⁸ As a result, early upright positioning is not universally accepted by stroke physicians.

Little is known in patients with acute stroke about the influence of different body positions on blood pressure (BP) or on real-time cerebral flow variables. Transcranial Doppler (TCD) may be used to assess CBF velocity (CBFV) at the stroke unit bedside. Nursing patients 'heads down' has been advocated, based on a 20% augmentation of CBFV when comparing supine with 30° head elevation in 20 patients severely affected with stroke.⁹ If such relatively minor changes in head position have such an impact, what would the consequences of more pronounced changes in position be in a representative sample of stroke unit patients?

In this study, we sought to assess the impact of the various head positions on CBFV comparing (1) hemispheres affected by stroke and unaffected hemispheres and (2) affected hemispheres and healthy controls. Any effect of upright positioning on neurological status and outcome was explored.

METHODS

Study design and patients

This observational study was conducted in the stroke unit of a teaching hospital between August 2008 and May 2011. Patients aged over 18 years with a first ever ischaemic stroke in the middle cerebral artery (MCA) territory were considered eligible. The measurements were taken on the stroke unit and mainly because of practical reasons (installation on stroke unit, informed consent approval and no interference with urgent clinical studies or consultations) performed in the subacute phase. Our protocol dedicated that the measurements had to be completed between 6 and 16 h after stroke symptom onset, mimicking the period of early in bed mobilisation initiation during daytime. Patients had to be conscious, cooperative, haemodynamically stable without fever and display flow in both M1 segments of the MCA. Moreover, patients deemed to be too severely affected by the stroke or too severely ill to be positioned upright (clinical/nursing judgement) were not included in this study. Brain CT, ECG, bilateral duplex carotid ultrasound and laboratory investigations were

performed. Thrombolytic therapy was administered according to (inter)national guidelines. Use of any medication, especially antiarrhythmics and antihypertensive medication, was registered. The National Institutes of Health Stroke Scale score (NIHSS) was used to determine neurological deficit. The stroke subtype was classified using the Oxfordshire Community Stroke Project classification,¹⁰ and stroke aetiology according to Trial of Org 10172 in Acute Stroke Treatment (TOAST) definitions.¹¹ The modified Rankin Scale (mRS) score at 3 months was determined by telephone interview by a certified assessor blinded to study data. Twenty healthy controls were recruited at random from (retired) work associates and their medical histories were reviewed. All experiments complied with the tenets of the Declaration of Helsinki and the study protocol was reviewed and approved by our institutional review board. Study participants provided written informed consent. Next of kin gave consent when participants were unable to provide it.

Sample size calculation

In the literature, SD of the intraindividual supine to sitting MCA CBFV difference is estimated to be around $\pm 5\%$ for elderly hospitalised patients¹² and $\pm 20\%$ for patients with severe stroke going from the 30° to 0° position.⁹ To detect a difference in mean CBFV of more than 6 cm/s (10–20%) between remaining supine and sitting upright in bed with a probability (power) of 90% when testing at a two-sided significance level of 5%, a sample size of 45 participants was required.

Procedures

BP was measured with a finger plethysmograph (Finometer-Pro, Finapres Medical Systems, Amsterdam, the Netherlands). The cuff was applied to the middle finger of the non-dominant or paretic hand placed at the heart level with the arm across the chest and supported by a sling. TCD was performed by the same investigator (MA) over the temporal bone window on both sides. MCA wave forms were identified at a depth range of 40–60 mm and a stable forward waveform with good intensity was selected for monitoring (Nicolet Pioneer TC8080, Carefusion Corporation, San Diego, California, USA). Patients with permanent MCA occlusion on TCD were excluded. TCD probes (2 MHz) were mounted by a head frame to ensure a constant angle of insonation throughout the positioning procedure. Bifrontal near-infrared spectroscopy (NIRS) recordings were added from 2010 onwards for regional cerebral oxygenation (rSO₂, %) calculation. The INVOS 5100C NIRS device (Somanetics Corp, Detroit, Michigan, USA) has a sampling frequency of 0.2 Hz. Adhesive optodes were placed on each side of the forehead. End-tidal CO₂ (EtCO₂, kPa) was measured by a mask using an infrared CO₂ analyser (Capnomac Ultima, GE Healthcare, Chalfont St Giles, UK) as an approximation of partial arterial CO₂ pressure.

Signals

The signals of the BP and EtCO₂ waveforms, spectral envelope of MCA CBFV, and rSO₂ as well as the position marker signals were sampled at 250 Hz and synchronously captured using an A/D converter. Signals were recorded with LabVIEW software (LabVIEW V.7.1, National Instruments, Austin, Texas, USA) for online inspection and were stored for postprocessing. The MCA CBFV_{mean} (cm/s) was calculated by using beat-to-beat mean values that were obtained by triggering off the ascending slope of the BP curve. Mean arterial pressure (MAP) at the MCA level (MAP_{mca}, mm Hg) was automatically calculated from MAP measured at the heart level and the vertical finger-to-TCD probe distance. In the upright and Trendelenburg positions, great care was taken to secure the hand with the finger plethysmograph cuff at the heart level (approaching the right atrium level). The beginning and ending time points of the postural transitions were marked, but these data periods (± 30 s) were not used for analysis.

Predefined positioning protocol

After admission to the stroke unit, patients were instrumented. Participants were requested to abstain from moving and talking in order to minimise artefacts and activity-related CBFV changes. Great care was taken to keep the head in the neutral midline position (without shifting of the headband) and the BP cuff at the heart level during positioning. A test run was performed. After 5 min of supine rest (baseline (eg, first) supine), the participants were passively tilted by changing the head of the bed end angles to the 45° (half sitting), 70° (sitting) and -15° (Trendelenburg) positions for 3–5 min each. Patients were brought in the sitting position twice. Periods of tilts were interspaced with 3–5 min of supine rest to re-establish control values (supine position). Before and after establishing each position, the motor score of NIHSS (affected arm and leg) was evaluated. Control participants were recorded in a quiet, temperature-controlled research room using the same predefined positioning protocol.

Statistical analysis

We calculated a mean value for every position period. To compare responses in each position, percentage changes in the variables (MCA CBFV_{mean}, rSO₂, MAP_{mca}, heart rate (HR) and EtCO₂) were calculated as the differences between the positions after normalisation to the mean baseline (eg, first) supine value. Normalisation was carried out to be able to compare physiological responses between patients, healthy controls and hemispheres, where absolute individual values might differ from baseline. We averaged these percentage changes to obtain a mean percentage change for each of the four different positions: supine, half sitting, sitting and Trendelenburg. For control participants, the mean of the MCA CBFV_{mean} and rSO₂ signals was taken. To account for and to take advantage of the correlation between repeated measures

in the same individual, a multilevel model was used to estimate the effects of positioning and affected hemisphere on changes in the main variables (MCA CBFV_{mean} and rSO₂).¹³ The relation between MCA CBFV_{mean} and outcome (dichotomised as mRS 0–2 for favourable outcome) was analysed in a similar way. Statistical analysis was performed with SAS (V.9.2). Significant carotid stenosis, stroke subtype or aetiology (TOAST criteria), NIHSS, age and sex were used as potential confounding covariates in stroke group comparisons.¹⁴ Age, sex and history of hypertension, cardiac disease and diabetes were used as covariates for healthy control and stroke group comparisons. Bonferroni adjustments for multiple comparisons were used for all results presented.

RESULTS

In eight patients with subacute stroke, TCD or BP investigation was unsuccessful. Eventually, we enrolled 47 patients (table 1). The mean patient age was 62 ± 15 years and median baseline NIHSS was 7 points (IQR 5–14 points). At 3 months, 29 patients (62%) had a favourable outcome. NIRS recordings were performed in 32 patients (68%) and 16 controls (80%).

We compared the cerebral haemodynamic postural (MCA CBFV_{mean} and rSO₂) changes between positions, patients with stroke and healthy controls, and between affected and unaffected hemispheres. A complete overview of the comparisons is presented in online supplementary tables 1–3 (<http://bmjopen.bmj.com>). Figure 1 shows that cerebral haemodynamic changes were most pronounced when moving from the supine to the sitting position (70°) for patients (affected hemisphere) and healthy controls (figure 1). The results presented below therefore focus on changes from the supine to the sitting position. The accompanying systemic haemodynamic (MAP_{mca} and HR) and respiratory responses (EtCO₂) are shown in figure 2.

In the sitting position for the affected and unaffected hemispheres, MCA CBFV_{mean} decreased in 28 patients (60%) and rSO₂ decreased in 18 patients (55%). None of the univariate results presented below were different after adjustment for predefined covariates, except for the outcome analysis.

Positional CBFV changes on sitting in patients with stroke

The mean MCA CBFV_{mean} decreased significantly with movement to the sitting position, (mean difference -4%, 95% CI -6% to -3%; $p=0.005$) for the affected hemisphere. For the unaffected hemisphere, this effect was not significant (mean difference -2%, 95% CI -5% to 0%; $p=1.0$).

Positional CBFV changes on sitting comparing patients with stroke and healthy controls

The mean MCA CBFV_{mean} changes on sitting were comparable between healthy controls and affected hemispheres of

Table 1 Patients' demographics and clinical characteristics

Variables	Patients with stroke (n=47)	Healthy controls (n=20)
Mean age (SD), years	62 (15)	52 (20)
Male	32 (70)	11 (55)
Affected right hemisphere	24 (52)	—
Median NIHSS score on admission (IQR)	7 (5–14)	—
History of hypertension	21 (46)	7 (35)
History of cardiac disease	17 (37)	0 (0)
Significant carotid stenosis	8 (17)	Unknown
Type infarct		
Atherosclerotic	15 (33)	
Cardiac	10 (22)	
Small vessel	14 (30)	
Dissection	2 (4)	
Unknown	5 (11)	
Oxford classification		
Total anterior circulation infarction	9 (20)	
Partial anterior circulation infarction	19 (41)	
Lacunar circulation infarction	18 (40)	
Posterior circulation infarction	0 (0)	
History of diabetes mellitus	8 (17)	0 (0)
Antihypertensive medication on admission	25 (54)	6 (30)
Intravenous thrombolysis	20 (44)	—
Mean arterial pressure (SD), mm Hg	84 (21)	78 (13)
Mean heart rate (SD), bpm	77 (18)	64 (7)

Values are absolute numbers (%), unless otherwise stated.
 NIHSS, National Institutes of Health Stroke Scale.

patients with stroke (mean difference -4% for the affected hemisphere (95% CI -11% to 3% ; $p=1.0$) as well as unaffected hemispheres (mean difference -5% , 95% CI -10% to 1% ; $p=0.18$)).

Positional CBFV changes on sitting comparing hemispheres in patients with stroke

The mean MCA CBFV_{mean} was not different between the affected and the unaffected hemispheres in the

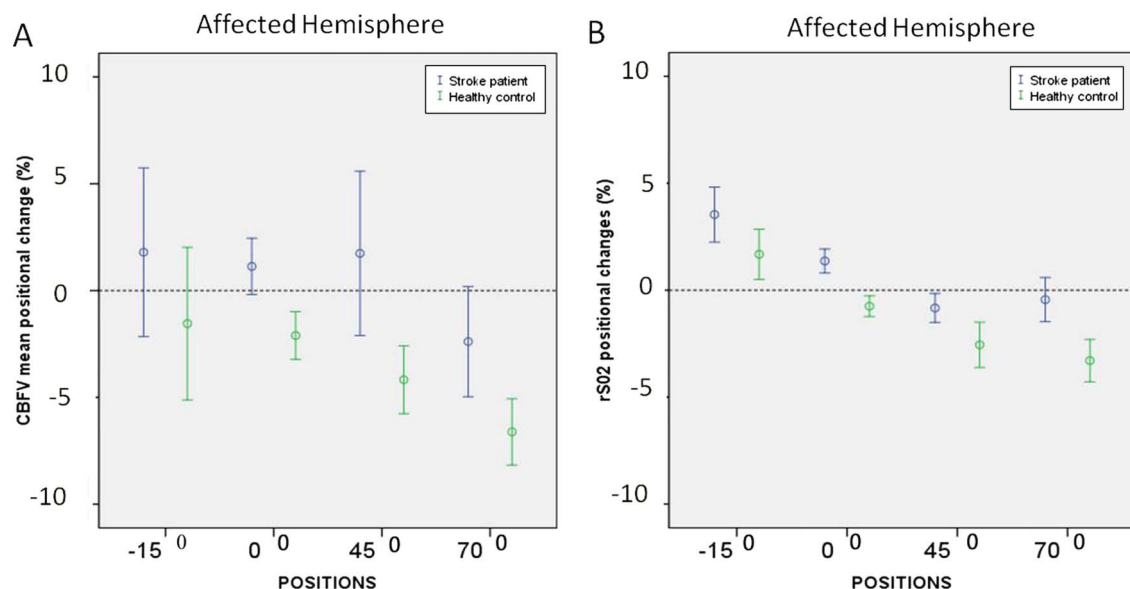


Figure 1 Cerebral haemodynamic changes in different body positions in patients with subacute stroke and healthy controls. Image showing changes of MCA CBFV_{mean} (A) and rSO₂ (B) in different positions following subacute stroke in patients (affected hemisphere) and healthy controls. All data were normalised to the baseline (eg, first) supine position and displayed in percentages (%). Thirty-two patients had near-infrared spectroscopy recordings. CBFV, cerebral blood flow velocity; MCA, middle cerebral artery; rSO₂, regional cerebral oxygen saturation.

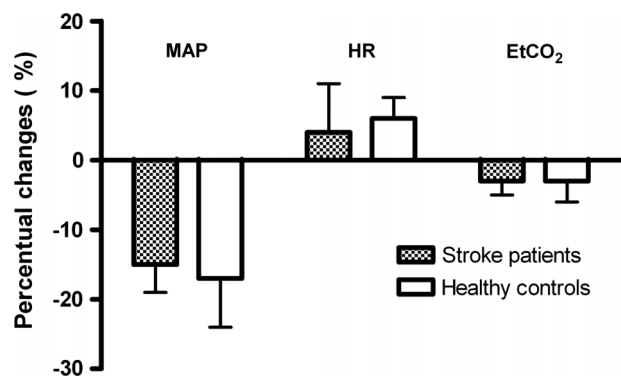


Figure 2 Systemic haemodynamic changes with going to the upright position in patients with subacute stroke and healthy controls. Image showing the changes of mean arterial pressure at the middle cerebral artery level (MAP_{mca}), heart rate (HR) and end-tidal CO_2 ($EtCO_2$) with going to the upright position (70°) in patients with subacute stroke and healthy controls. All data were normalised to the baseline (eg, first supine position and displayed in percentages (%).

sitting position (mean difference -2% , 95% CI -5% to 2% ; $p=1.0$).

Positional changes of rSO_2 , MAP_{mca} , HR and $EtCO_2$ on sitting

The mean rSO_2 values decreased significantly with movement to the sitting position for the affected hemisphere (mean difference -2% , 95% CI -3% to -1% ; $p=0.006$ and unaffected hemisphere (mean difference -2% , 95% CI -3% to -1% ; $p=0.009$). In agreement with the $MCA\ CBFV_{mean}$ results, the mean rSO_2 postural responses were not significantly different between both hemispheres and between patients and healthy controls. The $CBFV_{mean}$ and rSO_2 changes were positively correlated for the affected hemisphere ($r=0.33$, $p=0.01$).

In patients with stroke, MAP_{mca} decreased on average by 15% in the sitting position (95% CI -19% to -10% ; $p<0.001$). This was similar to controls (17% , 95% CI -24% to -11% ; $p<0.001$; figure 2). MAP_{mca} increased slightly during the course of the experiment for patients (mean 5% , 95% CI 2% to 7%) and healthy controls (mean 2% , 95% CI -1% to 6%). The postural responses were accompanied by only minor HR changes. $EtCO_2$ decreased on average by 3% (95% CI -5% to -1% ; $p<0.001$) for patients in the sitting position.

Positional changes, neurological status and relation with outcome in patients with stroke

We observed no neurological worsening or improvement during the postural changes in any of the patients. There were no significant differences between the mean $MCA\ CBFV_{mean}$ changes on upright and Trendelenburg positioning in patients with unfavourable compared to favourable outcome after correction for confounding covariates (mean difference on sitting, for the affected hemisphere: -5% , 95% CI -14% to 5% ; $p=1.0$).

DISCUSSION

Our study showed that following an ischaemic stroke in the MCA territory, early upright positioning in bed resulted in minor average changes in $CBFV$ only in the subacute phase. The postural responses were not significantly different between the affected and unaffected hemispheres, and between patients and healthy controls. Passive upright positioning was not accompanied by worsening of the (motor) stroke severity score. No improvement was noticed in the supine and Trendelenburg positions. The relative postural haemodynamic and $EtCO_2$ changes were largely comparable between patients and controls. This study found no indications that upright positioning compromises flow or (frontal) oxygenation significantly in the affected MCA territory in mild to moderately affected patients with acute stroke. No association was found between flow changes and functional outcome. Our results suggest effective cerebral and/or systemic haemodynamic adjustments to overcome gravitational force during postural changes in the subacute phase of mild to moderate stroke. The potential of the Trendelenburg position to augment CBF seems to be limited and may not outweigh concurrent cardiopulmonary complications.

Although TCD measurements cannot be used to calculate CBF, the observed relative $CBFV$ changes can be considered proportionate to CBF changes as long as the angle of insonation and the MCA diameter remain constant during the brief test interval. The MCA diameter is thought to remain constant over a ± 30 mm Hg range of BP and ± 2 kPa of CO_2 .¹⁵ This requirement was fulfilled in our study. TCD is prone to several errors because the measurements are operator dependent and patient dependent. Even small probe position changes may alter the $CBFV$ readings. However, frontal cerebral oxygenation (rSO_2) followed $CBFV$ changes in size and direction in the affected hemisphere, which further strengthens our findings. Assuming a constant arterial O_2 content and cerebral metabolic rate during the measurement, rSO_2 is predominantly a function of local CBF.^{16 17} Furthermore, TCD monitoring is the only non-invasive method to monitor CBF changes continuously at the bedside. It is generally accepted that CBF changes are closely reflected by TCD readings.¹⁸

Patients in our study predominantly had mild to moderate strokes. Patients with MCA occlusion were excluded. It was left to clinical and nursing judgement whether patients were deemed too severely affected by the stroke or too severely ill to be positioned upright for this study. Although NIHSS was not found to be a confounder for postural $CBFV$ changes, too few patients with severe stroke (NIHSS >16) were included to be able to generalise our findings to patients with severe strokes. Furthermore, fully ambulatory patients may have been less likely to be included. The practical implication of the latter drawback in generalisation is probably limited. When mild to moderate strokes are unlikely to be affected by upright positioning, very mild strokes are even less likely to be affected.

Three studies have investigated the postural effects on CBFV in acute stroke.^{9 19 20} Wojner *et al* found on average a 20% CBFV increase in the supine position in 20 patients with an acute MCA occlusion. However, a wide range of individual flow improvements was present (5–96% from baseline), with no data from the unaffected hemisphere being available.⁹ A small study with eight participants found important postural flow changes in four severely affected patients with incomplete MCA recanalisation on 24 h follow-up imaging (affected hemisphere).¹⁹ A more than 15% higher MCA flow was found in the supine position in 18 patients with large stroke with intracranial pressure (ICP) monitoring in the intensive care unit. This effect was only present in the affected hemisphere.²⁰ Different results were obtained in a single-photon emission CT (SPECT) stroke study. Seven of eight patients with subacute stroke had a clear visual improvement in SPECT perfusion to the affected frontal, temporal or parietal lobes in the semirecumbent (30–45°), compared to the supine position. No differences were found in BP in both positions. Unfortunately, no demographics, clinical effect or respiratory changes of the more upright position were reported.²¹

Our study did not show relevant postural changes despite the larger position changes (70° instead of 30°). This is probably largely due to our predominant focus on mild to moderately affected patients with stroke. Also, in our study, changes in all variables were related to the individual baseline supine level (by normalisation), reducing the effects of large variations of measured parameters and difficulties with standardisation of the used techniques (TCD, NIRS and BP). In none of the four studies were flow data recorded simultaneously with NIRS and EtCO₂. In the study of Hargroves *et al*, cerebral oxygenation as measured by NIRS in the first seven days after stroke was lowest in the upright position and highest in the supine position (difference estimated around 2–6%). These changes were also present in the unaffected hemisphere, although on visual inspection they were less marked.²² Furthermore, there might be an effect of anaesthesia, vasopressors, hypothermia and decompressive craniectomy on cerebral vasoregulation used in the intensive care stroke positioning study.²³ Our results are in line with patients with subarachnoid haemorrhage in whom CBFV remained constant in the supine and 45° patient body positions, regardless of the development of vasospasm.²⁴

It is widely believed that a sudden BP reduction can affect CBF and render elderly participants or patients with cardiovascular disease vulnerable to orthostatic symptoms.²⁵ The sitting position might be dangerous, especially with globally impaired cerebral autoregulation, because of the susceptibility of the penumbral regions to local hypoperfusion in the acute and subacute phases. However, our physiological results indicate that sitting is probably well tolerated in mild to moderately affected patients with stroke. In 2008, the 'A Very Early Rehabilitation trial' showed that very early out of

bed mobilisation was safe and feasible in patients with stroke.²⁶ This may be due to the counterbalancing effects of BP elevation, ICP reduction and increased venous return to stabilise cerebral perfusion pressure in the subacute phase. There is evidence for increased sympathetic activation in patients with acute stroke.^{27–29} Replication of this study should be considered in studying the (hyper)acute phase, longer sitting periods and using more sophisticated and direct methods that measure CBF regionally and estimate cerebral oxygen metabolism.³⁰ Inclusion of more severely disabled patients (NIHSS around 10) would be preferable to being able to compare upright cerebral haemodynamics with the results from the ongoing 'early mobilisation' trials.²⁶ Further research should also include more detailed (follow-up) information from extracranial and intracranial vessels pathology (using validated radiological or ultrasound criteria), recanalisation (proximal and distal), penumbra configuration and/or the presence of collaterals.^{31–34} Follow-up studies several days after stroke may be warranted because local and systemic compensatory mechanisms may become exhausted, but around that time influence on penumbral salvage may be limited.³⁵ Our results do not hold for sitting to the standing mobilisation, which may provoke different (probably larger) haemodynamic responses.³⁶

In the paucity of results for the early out of bed mobilisation stroke trials,³⁷ we suggest continuation of early upright positioning in patients with stroke in the stroke unit in the subacute phase and propose that continuous CBF measurements during early activity out of bed (standing) could be helpful in assuring safe and early mobilisation in the stroke unit.³⁶

CONCLUSION

In conclusion, upright positioning in the stroke unit of mildly to moderately affected patients with stroke does not give significant reduction of CBF and (frontal) cerebral oxygenation surrogates during the subacute phase on the stroke unit. Supine or Trendelenburg positioning seems not to augment MCA CBFV_{mean} or improve neurological status.

Contributors MJA, JWE and PV contributed to design and conceptualisation of the study, interpretation of the data, and drafting and revision of the manuscript. RS and JDK contributed to the interpretation of the data and revision of the manuscript. BK was responsible for the revision of the manuscript.

Funding MJA is supported by the Netherlands Organisation for Health Research and Development.

Competing interests None.

Patient consent Obtained.

Ethics approval Medical Ethical Committee, University Medical Center Groningen, Groningen, the Netherlands.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement The authors are willing to share the data, as well as the additional unpublished data of follow-up studies (day 3 after stroke) and dynamic autoregulation results.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 3.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/3.0/>

REFERENCES

1. Markus HS. Cerebral perfusion and stroke. *J Neurol Neurosurg Psychiatry* 2004;75:353–61.
2. Diserens K, Michel P, Bogousslavsky J. Early mobilisation after stroke: review of the literature. *Cerebrovasc Dis* 2006;22:183–90.
3. Bernhardt J, Indredavik B, Dewey H, *et al.* Mobilisation 'in bed' is not mobilisation. *Cerebrovasc Dis* 2007;24:157–8.
4. Hill K. Australian Clinical Guidelines for Acute Stroke Management 2007. *Int J Stroke* 2008;3:120–9.
5. Tyson SF, Nightingale P. The effects of position on oxygen saturation in acute stroke: a systematic review. *Clin Rehabil* 2004;18:863–71.
6. McArthur KS, Quinn TJ, Higgins P, *et al.* Post-acute care and secondary prevention after ischaemic stroke. *BMJ* 2011;342:d2083.
7. Cumming TB, Thrift AG, Collier JM, *et al.* Very early mobilization after stroke fast-tracks return to walking: further results from the phase II AVERT randomized controlled trial. *Stroke* 2011;42:153–8.
8. Skarin M, Bernhardt J, Sjöholm A, *et al.* 'Better wear out sheets than shoes': a survey of 202 stroke professionals' early mobilisation practices and concerns. *Int J Stroke* 2011;6:10–15.
9. Wojner-Alexander AW, Garami Z, Chernyshev OY, *et al.* Heads down: flat positioning improves blood flow velocity in acute ischemic stroke. *Neurology* 2005;64:1354–7.
10. Bamford J, Sandercock P, Jones L, *et al.* The natural history of lacunar infarction: the Oxfordshire Community Stroke Project. *Stroke* 1987;18:545–51.
11. Kolominsky-Rabas PL, Weber M, Gefeller O, *et al.* Epidemiology of ischemic stroke subtypes according to TOAST criteria: incidence, recurrence, and long-term survival in ischemic stroke subtypes: a population-based study. *Stroke* 2001;32:2735–40.
12. Poli L, Bo M, Secreto P, *et al.* Age-related transcranial Doppler findings in the evaluation of cerebral circulation during postprandial and postural tests. *Cerebrovasc Dis* 1999;9:98–101.
13. Littell RC, Milliken GA, Stroup WW, *et al.* *SAS system for mixed models*. North Carolina: SAS Institute Inc, 1996.
14. Immink RV, van Montfrans GA, Stam J, *et al.* Dynamic cerebral autoregulation in acute lacunar and middle cerebral artery territory ischemic stroke. *Stroke* 2005;36:2595–600.
15. Giller CA, Bowman G, Dyer H, *et al.* Cerebral arterial diameters during changes in blood pressure and carbon dioxide during craniotomy. *Neurosurgery* 1993;32:737–41.
16. Bucher HU, Edwards AD, Lipp AE, *et al.* Comparison between near infrared spectroscopy and ¹³³Xenon clearance for estimation of cerebral blood flow in critically ill preterm infants. *Pediatr Res* 1993;33:56–60.
17. Rasmussen P, Dawson EA, Nybo L, *et al.* Capillary-oxygenation-level-dependent near-infrared spectrometry in frontal lobe of humans. *J Cereb Blood Flow Metab* 2007;27:1082–93.
18. Newell DW, Aaslid R, Lam A, *et al.* Comparison of flow and velocity during dynamic autoregulation testing in humans. *Stroke* 1994;25:793–7.
19. Hunter AJ, Snodgrass SJ, Quain D, *et al.* HOBEO (Head-of-Bed Optimization of Elevation) Study: association of higher angle with reduced cerebral blood flow velocity in acute ischemic stroke. *Phys Ther* 2011;91:1503–12.
20. Schwarz S, Georgiadis D, Aschoff A, *et al.* Effects of body position on intracranial pressure and cerebral perfusion in patients with large hemispheric stroke. *Stroke* 2002;33:497–501.
21. Jack CI, Lee T, Moyle P, *et al.* The importance of posture in the early stages of stroke: its influence on cerebral perfusion. *Age Ageing* 2001;30:428.
22. Hargroves D, Tallis R, Pomeroy V, *et al.* The influence of positioning upon cerebral oxygenation after acute stroke: a pilot study. *Age Ageing* 2008;37:581–5.
23. Dagal A, Lam AM. Cerebral autoregulation and anesthesia. *Curr Opin Anaesthesiol* 2009;22:547–52.
24. Zhang Y, Rabinstein AA. Lower head of the bed position does not change blood flow velocity in subarachnoid hemorrhage. *Neurocrit Care* 2011;14:73–6.
25. Lipsitz LA. Orthostatic hypotension in the elderly. *N Engl J Med* 1989;321:952–7.
26. Bernhardt J, Dewey H, Thrift A, *et al.* A very early rehabilitation trial for stroke (AVERT): phase II safety and feasibility. *Stroke* 2008;39:390–6.
27. Qureshi AI. Acute hypertensive response in patients with stroke: pathophysiology and management. *Circulation* 2008;118:176–87.
28. Korpelainen JT, Sotaniemi KA, Suominen K, *et al.* Cardiovascular autonomic reflexes in brain infarction. *Stroke* 1994;25:787–92.
29. Chamorro A, Amaro S, Vargas M, *et al.* Catecholamines, infection, and death in acute ischemic stroke. *J Neurol Sci* 2007;252:29–35.
30. Ouchi Y, Yoshikawa E, Kanno T, *et al.* Orthostatic posture affects brain hemodynamics and metabolism in cerebrovascular disease patients with and without coronary artery disease: a positron emission tomography study. *Neuroimage* 2005;24:70–81.
31. Tsvigoulis G, Ribo M, Rubiera M, *et al.* Real-time validation of transcranial Doppler criteria in assessing recanalization during intra-arterial procedures for acute ischemic stroke: an international, multicenter study. *Stroke* 2013;44:394–400.
32. Bang OY, Saver JL, Buck BH, *et al.* Impact of collateral flow on tissue fate in acute ischaemic stroke. *J Neurol Neurosurg Psychiatry* 2008;79:625–9.
33. Wheeler HM, Mlynash M, Inoue M, *et al.* Early diffusion-weighted imaging and perfusion-weighted imaging lesion volumes forecast final infarct size in DEFUSE 2. *Stroke* 2013;44:681–5.
34. Tomsick T. TIMI, TIBI, TICI: I came, I saw, I got confused. *AJNR Am J Neuroradiol* 2007;28:382–4.
35. Reinhard M, Wihler C, Roth M, *et al.* Cerebral autoregulation dynamics in acute ischemic stroke after rtPA thrombolysis. *Cerebrovasc Dis* 2008;26:147–55.
36. Kim YS, Bogert LW, Immink RV, *et al.* Effects of aging on the cerebrovascular orthostatic response. *Neurobiol Aging* 2011;32:344–53.
37. Bernhardt J, Dewey H, Collier J, *et al.* A Very Early Rehabilitation Trial (AVERT). *Int J Stroke* 2006;1:169–71.

BMJ Open

Cerebral blood flow velocity changes during upright positioning in bed after acute stroke: an observational study

Marcel J Aries, Jan Willem Elting, Roy Stewart, Jacques De Keyser, Berry Kremer and Patrick Vroomen

BMJ Open 2013 3:

doi: 10.1136/bmjopen-2013-002960

Updated information and services can be found at:
<http://bmjopen.bmj.com/content/3/8/e002960>

These include:

Supplementary Material

Supplementary material can be found at:
<http://bmjopen.bmj.com/content/suppl/2013/08/13/bmjopen-2013-002960.DC1>

References

This article cites 36 articles, 16 of which you can access for free at:
<http://bmjopen.bmj.com/content/3/8/e002960#BIBL>

Open Access

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 3.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/3.0/>

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections

Articles on similar topics can be found in the following collections

[Neurology](#) (423)
[Rehabilitation medicine](#) (312)

Notes

To request permissions go to:
<http://group.bmj.com/group/rights-licensing/permissions>

To order reprints go to:
<http://journals.bmj.com/cgi/reprintform>

To subscribe to BMJ go to:
<http://group.bmj.com/subscribe/>